

SUBCHAPTER E—PESTICIDE PROGRAMS

PART 150—GENERAL

AUTHORITY: Reorganization Plan No. 3 of 1970 (5 U.S.C. App.).

§ 150.17 Addresses for applications and correspondence.

The official addresses for all submissions directed to the Office of Pesticide Programs (OPP) of the Environmental Protection Agency are as follows:

(a) *United States Postal Service mailing address.* Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave., NW., Washington DC 20460-0001.

(b) *Hand/courier delivery address.* Office of Pesticide Programs, Environmental Protection Agency, 2777 S. Crystal Dr., Arlington, VA 22202-4501.

(c) *OPP Regulatory Public Docket address.* OPP Regulatory Public Docket is physically located in Rm. S-4400, One Potomac Yard (South Bldg.), 2777 S. Crystal Dr., Arlington, VA 22202-4501. This is not a mailing address.

[71 FR 35545, June 21, 2006]

PART 151 [RESERVED]

PART 152—PESTICIDE REGISTRATION AND CLASSIFICATION PROCEDURES

Subpart A—General Provisions

Sec.

152.1 Scope.

152.3 Definitions.

152.5 Pests.

152.6 Substances excluded from regulation by FIFRA.

152.8 Products that are not pesticides because they are not for use against pests.

152.10 Products that are not pesticides because they are not deemed to be used for a pesticidal effect.

152.15 Pesticide products required to be registered.

Subpart B—Exemptions

152.20 Exemptions for pesticides regulated by another Federal agency.

152.25 Exemptions for pesticides of a character not requiring FIFRA regulation.

152.30 Pesticides that may be transferred, sold, or distributed without registration.

Subpart C—Registration Procedures

152.40 Who may apply.

152.42 Application for new registration.

152.43 Alternate formulations.

152.44 Application for amended registration.

152.46 Notification and non-notification changes to registrations.

152.50 Contents of application.

152.55 Where to send applications and correspondence.

Subpart D [Reserved]

Subpart E—Procedures To Ensure Protection of Data Submitters' Rights

152.80 General.

152.81 Applicability.

152.83 Definitions.

152.84 When materials must be submitted to the Agency.

152.85 Formulators' exemption.

152.86 The cite-all method.

152.90 The selective method.

152.91 Waiver of a data requirement.

152.92 Submission of a new valid study.

152.93 Citation of a previously submitted valid study.

152.94 Citation of a public literature study or study generated at government expense.

152.95 Citation of all studies in the Agency's files pertinent to a specific data requirement.

152.96 Documentation of a data gap.

152.97 Rights and obligations of data submitters.

152.98 Procedures for transfer of exclusive use or compensation rights to another person.

152.99 Petitions to cancel registration.

Subpart F—Agency Review of Applications

152.100 Scope.

152.102 Publication.

152.104 Completeness of applications.

152.105 Incomplete applications.

152.107 Review of data.

152.108 Review of labeling.

152.110 Time for Agency review.

152.111 Choice of standards for review of applications.

152.112 Approval of registration under FIFRA sec. 3(c)(5).

152.113 Approval of registration under FIFRA sec. 3(c)(7)—Products that do not contain a new active ingredient.